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I. AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (currently amended) A method for treating a person suffering from retinal

edema or non-proliferative diabetic retinopathy which comprises, administering an effective

amount of a composition comprising a glucocorticoid and anecortave acetate, wherein said

composition is free of classical preservativespreservative-free.

2. (cancel)

3. (previously presented) The method of claim 1, wherein the glucocorticoid is

selected from the group consisting of dexamethasone, fluoromethalone, medrysone,

betamethasone, triamcinolone, triamcinolone acetonide, prednisone, prednisolone,

hydrocortisone, rimexolone, and pharmaceutically acceptable salts thereof.

4. (previously presented) The method of claim 1, wherein the glucocorticoid is

selected from the group consisting of prednicarbate, deflazacort, halomethasone, tixocortol,

prednylidene (21-diethylaminoacetate), prednival, paramethasone, methylprednisolone,

meprednisone, mazipredone, isoflupredone, halopredone acetate, halcinonide, formocortal,

flurandrenolide, fluprednisolone, fluprednidien acetate, fluperolone acetate, fluocortolone,

fluocortin buytl, fluocinonide, fluocinolone acetonide, flumisolide, flumethasone, fludrocortisone,

fluclorinide, enoxolone, difluprednate, diflucortolone, diflorasone diacetate, desoximetasone

(desoxymethasone), desonide, descinolone, cortivazol, corticosterone, cortisone, cloprednol,

clocortolone, clobetasone, clobetasol, chloroprednisone, cafestol, budesonide, beclomethasone,

amcinonide, allopregnane acetonide, alclometasone, 21-acetoxypregnenolone, tralonide,

diflorasone acetate, deacycortivazol, RU-26988, budesonide, and deacylcortivazol oxetanone.

5. (previously presented) The method of claim 3, wherein the glucocorticoid is

triamcinolone acetonide.

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6. (previously presented) The method of claim 5, wherein the concentration of

triamcinolone acetonide in the composition is from 0.4% to 4.0% w/v.

7. (currently amended) The method of claim $2\underline{3}$, wherein the glucocorticoid is

rimexolone.

8. (previously presented) The method of claim 7, wherein the concentration of

rimexolone in the composition is from 0.1% to 4.0% w/v.

9. (currently amended) The method of claim $2\underline{3}$, wherein the glucocorticoid is

prednisolone acetate.

10. (previously presented) The method of claim 9, wherein the concentration of

prednisolone acetate in the composition is from 0.1% to 2.0% w/v.

11. (currently amended) The method of claim $\frac{23}{2}$, wherein the glucocorticoid is

fluoromethalone acetate.

12. (previously presented) The method of claim 11, wherein the concentration of

fluoromethalone acetate in the composition is from 0.1% to 1.0% w/v.

13. (currently amended) The method of claim 6, wherein the concentration of

anecortave acetate in the composition is from 0.1% to 6% about 3% w/v and the concentration of

triamcinolone acetonide in the composition is from 0.5% to $\frac{2.0\%4.0\%}{4.0\%}$ w/v.

14. (cancelled)

15. (cancelled)

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16. (currently amended) The method of claim 23, wherein the composition is delivered by a method selected from the group consisting of intravitreal injection, posterior juxtascleral delivery, subconjunctival injection, and implanted device.

- 17. (previously presented) The method of claim 16, wherein the composition is delivered by posterior juxtascleral injection.
- 18. (previously presented) The method of claim 16, wherein the composition is delivered via an implanted device.